The listing of claims presented below replaces all prior versions and listings of claims in the application.

## IN THE CLAIMS

Claims 1-51 (cancel)

52. (Currently Amended) A biochip comprising an array of gel cells formed on a substrate by copolymerization of composition K, wherein,

$$K = aA + bB + cC + dD + eE$$
 wherein

A is a monomer based on derivatives of acrylic and methacrylic acids;

**B** is a water soluble cross-linking agent;

C is a biological modified macromolecule bearing an unsaturated group;

**D** is a water soluble compound as a medium component for performing a copolymerization;

E is water, and

a, b, c, d, and e are percentages (X) of each ingredient in the composition wherein for solids X is m/v×100%; and for liquids X is v/v×100% wherein the total content of monomer and cross-linking agent is in a range from 3 to 40% (3 ≤ (a+b) ≤40%), and a monomer to cross-linking agent ratio being within a range of 97:3 to 60:40 and percentages of C, D, and E ingredients being within a range of 0.0001% ≤ c ≤10%; 0%≤ d ≤90%; 5%≤ e≤95%; wherein the cross-linking agent B is one or more of is one or more of N,N'methylenebisacrylamide, N,N' ethylenebismethacrylamide, N,N'-(1,2-dihydroxyethylene)bisacrylamide, and polyethylene glycol diacrylateand wherein each cell may include an immobilized macromolecule.

53. (Previously Presented) The biochip according to claim 52 wherein said cells form a regular one- or two-dimensional structure (phase).

- 54. (Previously Presented) The biochip according to claim 54 wherein the composition K is applied to a substrate by using an automatic device equipped with one or more micro dispensers.
- 55. (Previously Presented) The biochip according to claim 54 wherein the micro dispensers are rod type.
- 56. (Previously Presented) The biochip according to claim 54 wherein the micro dispensers are contactless micro dispensers of jet type.
- 57. (Previously Presented) The biochip according to claim 54 wherein the micro dispensers form a regular structure.

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- 58. (Previously Presented) The biochip according to claim 52 wherein one or more substrates including applied droplets of polymerization mixture, during polymerization, are placed into a sealed container under oxygen free inert atmosphere with a controlled humidity.
- 59. (Previously Presented) The biochip according to claim 58 wherein said container is filled with N<sub>2</sub>, Ar, or CO<sub>2</sub> gas.
- 60. (Previously Presented) The biochip according to claim 59 wherein the gas is continuously or periodically added to the container.
- 61. (Previously presented) The biochip according to claim 52 wherein monomer A is one or more of acrylamide, methacrylamide, N[tris(hydroxymethyl)methyl]acrylamide, and 2-hydroxyethylmethacrylate.
  - 62. (Previously Presented) The biochip according to claim 52 wherein monomers are used separately or as a mixture.
  - 63. (Cancelled)
  - 64. (Previously Presented). The biochip according to claim 52 wherein the cross-linking agents are used separately or as a mixture.
  - 65. (Previously Presented) The biochip according to claim 52 wherein the modified biological macromolecule C is of formula (I):

wherein

OLIGO represents an oligonucleotide;

 $R^1,\,R^2$ , and  $R^3$  are different and are selected from H, alkyl  $C_1\text{-}C_6,\,Ph,\,$  and  $PhCH_2\text{-}$ ;

Z is  $(CH_2)_nCH(CH_2OH)CH_2OX$  where n is 1-6; or Z is  $(CH_2)_r$ -OX where r is 2-6;

X is a phosphodiester group binding an unsaturated moiety to 5'- and/or 3'- end of the oligonucleotide;

 $R^4$  represents H, or  $(CH_2)_rOH$  where r is 2-6; and Y is  $(p-C_6H_4)_t$  where t is 0-2.

66. (Withdrawn) The biochip according to claim 52 wherein the modified biological macromolecule C is of formula (II):

wherein

DNA represents a DNA fragment,

X is H or H<sub>2</sub>PO<sub>3</sub>, and Z represents -CO-Y-CR<sup>1</sup>=CR<sup>2</sup>R<sup>3</sup>

or

X is  $-CO-Y-CR^1=CR^2R^3$ , and Z is H or  $H_2PO_3$ ;

 $R^1$ ,  $R^2$ , and  $R^3$  are the same different and are selected from H, alkyl  $C_1$ - $C_6$ , Ph, and PhCH<sub>2</sub>-; and

Y represents  $(p-C_6H_4)_t$  where t is 0-2.

67. (Withdrawn) The biochip according to claim 52 wherein the modified biological macromolecule C is of formula (III);

wherein:

DNA represents a DNA fragment;

 $R^1$ ,  $R^2$ ,  $R^3$  are the same different and are selected from H, alkyl  $C_1$ - $C_6$ , Ph, and PhCH<sub>2</sub>-; and

Y is  $(p-C_6H_4)_t$  where t is 0-2.

68. (Previously Presented) The biochip according to claim 52 wherein the modified biological macromolecule C is of formula (IV):

wherein:

## DNA represents a DNA fragment;

 $R^1,\,R^2,\,$  and  $R^3$  are the same different and are selected from H, alkyl  $C_1\text{-}C_6,\,$  Ph, and PhCH2– ; and

Y is  $(p-C_6H_4)_t$  where t is 0-2;

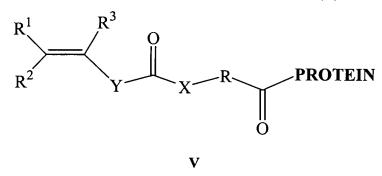
R<sup>4</sup> represents H, (CH<sub>2</sub>)<sub>r</sub>OH where r is 2-6; and

Z is (CH<sub>2</sub>)<sub>n</sub>CH(CH<sub>2</sub>OH)CH<sub>2</sub>OX where n is 1-6; or -(CH<sub>2</sub>)<sub>r</sub>-OX where r is 2-6;

and

X is a phosphodiester group binding an unsaturated moiety to 5'- and/or 3'- end of the DNA fragment.

69. (Withdrawn) The biochip according to claim 52 wherein the modified biological macromolecule C is a protein of formula (V):



wherein

 $R^1$ ,  $R^2$ , and  $R^3$  are the same different and are selected from H, alkyl  $C_1$ - $C_6$ , Ph, and PhCH<sub>2</sub>-;

X is NH, O, CH<sub>2</sub>, or S;

Y is  $(p-C_6H_4)_t$  where t is 0-2; and

R is  $(CH_2)_s$ , or  $(CH_2CH_2O)_s$ , where s is 1-20.

70. (Withdrawn) The biochip according to claim 52 wherein the modified biological macromolecule C is a protein of formula (VI):

wherein

 $R^1$ ,  $R^2$ , and  $R^3$  are the same different and are selected from H, alkyl  $C_1$ -  $C_6$ , Ph, and PhCH<sub>2</sub>-;

X is NH, O, S, or CH<sub>2</sub>;

Y is  $(p-C_6H_4)_t$ , where t is 0-2;

R is  $(CH_2)_s$ , or  $(CH_2CH_2O)_s$ , where s is 1-20;

W is NH, O, or CH<sub>2</sub>;

F is  $(CH_2)_x$ , where x is 1 or 2; and

Z is NH or S.

71. (Withdrawn) The biochip according to claim 52 wherein the modified biological macromolecule C is a protein of formula (VII):

wherein R represents (CH<sub>2</sub>)<sub>s</sub>, or (CH<sub>2</sub>CH<sub>2</sub>O)<sub>s</sub>, where s is 1-20.

72. (Previously Presented) The biochip according to claim 52 wherein D is a water soluble high-boiling organic compound.

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- 73. (Previously Presented) The biochip according to claim 72 where the water soluble high-boiling organic compound is *N,N*-dimethylformamide, dimethylsulfoxide or both.
- 74. (Previously Presented) The biochip according to claim 52 wherein use is made of a water soluble polyhydric compound as a component of the medium for performing the photo initiated polymerization.
- 75. (Previously Presented) The biochip according to claim 74 wherein the one or more water soluble polyhydric compound is selected from glycerol, sucrose and polyvinyl alcohol.
- 76. (Withdrawn) A method for performing PCR over the biochip according to claim 52 comprising the steps of:
  - a) adding amplification solution, forward (F) and reverse (R) primers of samples of nucleic acids under investigation; and
  - b) incubating the biochip under conditions of a thermocycling treatment providing a realization of PCR-amplification.
- 77. (Withdrawn) A method for performing the PCR over the biochip according to claim 52 comprising the steps of:
  - a) incubating isothermally the biochip with hybridization solution comprising the samples of nucleic acids under investigation to perform their hybridization with primers immobilized (synthetic oligonucleotides);
  - b) incubating isothermally the biochip, comprising the nucleic acids being hybridized with primers immobilized, in the amplification solution containing forward (F) and reverse (R) primers;
  - c) replacing the amplification solution out of biochip gel elements with hydrophobic liquid (mineral oil) which completely isolates biochip cells with each other, and

incubating the biochip under conditions of a thermocycling treatment providing a realization of PCR-amplification.

78. (New) A biochip comprising an array of gel cells formed on a substrate by copolymerization of composition K, wherein,

$$K = aA + bB + cC + dD + eE$$
 wherein

A is a monomer based on derivatives of acrylic and methacrylic acids;

B is a water soluble cross-linking agent;

C is a biological modified macromolecule bearing an unsaturated group;

**D** is a water soluble compound as a medium component for performing a copolymerization;

E is water, and

**a, b, c, d,** and **e** are percentages (X) of each ingredient in the composition wherein for solids X is m/v×100%; and for liquids X is v/v×100% wherein the total content of monomer and cross-linking agent is in a range from 3 to 40% (3  $\leq$  (**a+b**)  $\leq$ 40%), and a monomer to cross-linking agent ratio being within a range of 97:3 to 60:40 and percentages of **C**, **D**, and **E** ingredients being within a range of 0.0001%  $\leq$  **c**  $\leq$ 10%; 0%  $\leq$  **d**  $\leq$ 90%; 5%  $\leq$  **e** $\leq$ 95 and wherein each cell may include an immobilized macromolecule